





XX MO96C325-A1.  
 XX 04-FEB-1999.  
 XX 24-JUL-1998; 98WO-US15464.  
 XX 25-JUL-1997; 97US-0053097.  
 XX (UYBO-) UNIV BOSTON.  
 XX Goldstein RN;  
 XX WPI; 1999-142969/12.  
 XX Determining species of bacteria and fungi - useful for  
 XX distinguishing between bacterial/fungal species, and for determining  
 XX the identity of bacterial/fungal pathogens in biological samples  
 XX Disclosure: Fig 7 (60/67-63/67); 133pp; English.  
 XX This is the DNA sequence of the Escherichia coli strain MG1655  
 XX rrm operon (16S-spacer-23S-spacer-5S). Restriction sites for  
 XX enzymes cutting the operon 5 times or less have been determined.  
 XX E. coli rrmA-rrmH operon sequences are provided (see AAX4983-89).  
 XX Methods and compositions are described for determining the species  
 XX of an unknown bacterium or fungus in a sample. The method involves  
 XX isolating and digesting bacterial (or fungal) DNA encoding 16S and  
 XX 23S rRNA from a sample with restriction enzymes, detecting the  
 XX products, and comparing them to signature bands from a number of  
 XX bacteria. The method generates a species conserved set of RFLP  
 XX bands, unique for each species. These species-conserved sets  
 XX represent precise markers appropriate for inter-species  
 XX discriminatory purposes (i.e. to determine the species of a given,  
 XX unknown isolate e.g. in a clinical specimen). In contrast to  
 XX conventional ribotyping, the present invention utilizes the  
 XX ribosomal operon sequences which vary less than 3% (and more  
 XX preferably less than 2%) within a species and vary between species.  
 XX The method is useful for medical, food, agricultural and  
 XX environmental testing. It does not require sequencing of nucleic  
 XX acid from biological samples.  
 XX Sequence 5105 bp; 1334 A; 1133 C; 1565 G; 1065 T; 9 other:  
 XX  
 Query Watch 87.3%; Score 1270.8; DB 20; Length 5105;  
 Best Local Similarity 96.0%; Pred. No. 0;  
 Matches 1313; Conservative 0; Mismatches 54; Indels 1; Gaps 1;

QY 360 cgcctatgaagaagccttcgagttgttaaggaacttccacggaggaatgaagcttca 477  
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 QY 540 gcaagcgaagatgaagcgaaggaaggaaggaaggaaggaaggaaggaaggaaggaagga 569  
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 ID AAT29140 standard: DNA: 1542 bp.  
 XX















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RESULT 8
AAK24984
ID AAK24984 standard; DNA; 5098 bp.
AC AAK24984;
DT 05-JUL-1999 (first entry)
XX
DE E. coli MG1655 rmb operon (16S-spacer 23S spacer 58).
XX
KM Speciation, ribotyping, species discrimination, marker, rFLP,
KM restriction fragment length polymorphism; bacterium; fungus;
KM pathogen, rmb operon, 16S rRNA gene, 23S RNA gene, etc.
XX
OS Escherichia coli.
XX
FH Key Location/Qualifiers
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## RESULT 11

AA046119

AA046119 standard; tRNA; 1542 bp.

AA046119

22-FEB-1994 (first entry)

E. coli 16S rRNA fragment

Polymerase chain reaction (PCR) amplification of the 16S rRNA gene

Index pattern: electrophoresis; chromatography; ss

Synthetic

J005192147-A

61-AUG-1993

15-NOV-1991: 914P-0300882

15-NOV-1991: 914P-0300882

(KIRIN) KIRIN BEVERAGE KK

WP1: 1993-273464/55

Specification of DNA base sequence - by PCR amplification using

primers, cleavage resultant DNA with restriction enzyme and separ-

ation by electrophoresis or chromatography

Disclousure: Page 8-9; 20pp; Japanese

The sequences given in AA046119-20 are fragments of the 16S rRNA

sequence which was used to demonstrate the method of the

invention. Primers were used to amplify the target DNA which was

then cleaved. The fragments were separated by electrophoresis or

chromatography. This method can specify DNA sequences with high

reproducibility and can be applied to various industrial fields

such as chemistry, medical care, food stuffs and electronics.

Sequence 1512 bp; 489 A; 450 C; 486 G; 317 T; 0 other

Query Match 85.98; Score 1250; DB 14; Length 1542;

Fast Local Similarity 95.0%; Pred. No. 0;

Matches 1300; Conservative 0; Mismatches 67; Indels 1; Gaps 1;

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